## Amendments to the Claims

The following listing of claims will replace all prior versions and listings of claims in the application.

## Listing of Claims:

(Currently Amended) A compound having the formula I, or a pharmaceutically 1 acceptable salt thereof.

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wherein

Y is -NH-R<sup>2</sup> or a group of formula

$$-N \searrow_{R^b}^{R^a}$$

R1 is cycloalkyl or non-substituted alkyl.

R2 is cycloalkyl,

R3 is hydrogen, alkyl, halogen, hydroxy, alkoxy or amino,

or R2R2 is an alkylene bridging group,

Ra is hydrogen, alkyl, alkenyl, alkynyl, halogen, hydroxy, alkoxy, amino,

alkylamino, alkylsulfonyloxy, cyano, carboxy, ester or amido,

Rb is hydrogen, alkyl or halogen,

or RaRb is carbonyl,

R4 is hydrogen or alkyl,

R5 is cycloalkyl, arylalkyl or heterocycle-alkyl,

or NR<sup>4</sup>R<sup>5</sup> is a heterocycle, which may be substituted, containing only one heteroatom

which is a nitrogen atom or containing two heteroatoms wherein one is a nitrogen atom and the other is a non-oxidized sulfur atom.

with the proviso that when Y is NHR<sup>2</sup> and R<sup>3</sup>R<sup>3</sup> is an alkylene bridging group or when Y is a group of formula

$$-N \searrow_{R^b}^{R^a}$$

R1 is a cycloalkyl.

- 2. (Original) A compound according to claim 1 wherein Y is -NH-R<sup>2</sup>.
- 3. (Currently Amended) A compound according to claim 2 wherein

R1 is C3-7-cycloalkyl or non-substituted alkyl,

R<sup>2</sup> is C3-7-cycloalkyl,

R3 is hydrogen, C1-4-alkyl, halogen, hydroxy, alkoxy or amino,

or R2R2 is a C2-4-alkylene bridging group.

R4 is hydrogen or C1-4-alkyl,

R5 is C3-7-cycloalkyl, arylalkyl or heterocycle-alkyl,

- or NR<sup>4</sup>R<sup>5</sup> is a heterocycle, which may be substituted, containing only one heteroatom which is a nitrogen atom or containing two heteroatoms wherein one is a nitrogen atom and the other is a non-oxidized sulfur atom.
- (Currently Amended) A compound according to claim 2 wherein R<sup>1</sup> is C3-4-alkyl or C3-5-cycloalkyl, preferably-eyelopropyl, isopropyl, eyelobutyl, eyelopentyl, 2-methyl-eyelopropyl-or-eyelopropylmethyl.
- (Currently Amended) A compound according to claim 2 wherein
   R<sup>2</sup> is a C3-4-non-substituted cycloalkyl, or a cycloalkyl substituted by a C1-6-alkyl or an
   aryl, preferably cyclopropyl or cyclobutyl,
  - and/or R<sup>3</sup> is hydrogen, methyl, ethyl, a Cl atom, a F atom, a Br atom, amino or methoxy; o<del>r R<sup>2</sup> R<sup>3</sup> is an alkylene bridging group selected from ethylene, propylene and butylene</del>.
- (Currently Amended)A compound according to claim 2 wherein
   R<sup>4</sup> is hydrogen or C1-4-alkyl, preferably hydrogen or methyl,
   and/or R<sup>5</sup> is 2-(2-thienyl)ethyl, 2-furylmethyl, 2-thienylmethyl, 4-pyridinylmethyl,
   benzyl, 2-(methylsulfanyl)benzyl, 2,6-difluorobenzyl, 2-fluorobenzyl, 2-

nitrobenzyl, 3,5-bis(trifluoromethyl)benzyl, 3,5-difluorobenzyl, cyclohexyl, cyclohexyl, 4-methylcyclohexyl, or 2,2-diphenylethyl,

- or NR<sup>4</sup>R<sup>5</sup> is 1,3-thiazolidin-3-yl, 1-azepanyl, 1-azocanyl, 3,5-dimethyl-1-piperidinyl, 4(2-methoxyphenyl)-1-piperidinyl, 4-(hydroxy(diphenyl)methyl)-1-piperidinyl, 4(trifluoromethyl)-1-piperidinyl, 4,4-difluoro-1-piperidinyl, 4,4-dimethyl-1piperidinyl, 4-carbamoyl-1-piperidinyl, 4-benzyl-1-piperidinyl, 4-carboxy-1piperidinyl, 4-cyano-4-phenyl-1-piperidinyl, 4-ethoxycarbonyl-1-piperidinyl, 4ethyl-1-piperidinyl, 4-cthyl-4-methyl-1-piperidinyl, 4-hydroxy-1-piperidinyl, 4hydroxy-4-phenyl-1-piperidinyl, 4-hydroxymethyl-1-piperidinyl, 4-methyl-1piperidinyl, 4-methylene-1-piperidinyl, 4-oxo-1-piperidinyl, 3,6-dihydro-1(2H)pyridinyl, 3-azabicyclo[3.2.1]oct-3-yl, 4-thiomorpholinyl, 2-one-1-azepanyl, 3,4dihydro-2(1H)-isoquinolinyl, 1,4-dioxa-8-azaspiro[4.5]dec-8-yl, 1,3,3-trimethyl6-azabicyclo[3.2.1]oct-6-yl, octahydro-2(1H)-isoquinolinyl or 8-azaspiro[4.5]dec8-yl.
- (Previously Presented) A compound selected from 6-(1-azepanyl)-N,2-dicyclopropyl-5-methyl-4-pyrimidinamine; N,2-dicyclopropyl-6-(4,4-dimethyl-1-piperidinyl)-5-methyl-4-pyrimidin-amine; N,2-dicyclopropyl-5-methyl-6-(4-methyl-1-piperidinyl)-4-pyrimidinamine; 6-(3-azabicyclo[3.2.1]oct-3-yl)-N,2-dicyclopropyl-5-methyl-4-pyrimidinamine; N,2-dicyclo-propyl-5-methyl-6-(4-thiomorpholinyl)-4-pyrimidinamine; and pharmaceutically acceptable salts thereof.
- 8. (Original) A compound according to claim 1 wherein Y is a group of formula

$$-N \searrow_{R^b}^{R^a}$$

(Currently Amended) A compound according to claim 8 wherein NR <sup>4</sup>R<sup>5</sup> is a 5- to 9membered heterocycle, which may be substituted, containing only one heteroatom which
is a nitrogen atom or containing two heteroatoms wherein one is a nitrogen atom and the
other is a non-oxidized sulfur atom, preferably 1-azepanył.

10. (Original) A compound according to claim 9 wherein

R1 is C3-7-cycloalkyl,

R3 is hydrogen, C1-4-alkyl, halogen, hydroxy, alkoxy or amino,

R<sup>a</sup> is hydrogen, C1-4-alkyl, C2-6-alkenyl, C2-6-alkynyl, halogen, hydroxy, alkoxy, amino, alkylamino, alkylsulfonyloxy, cyano, carboxy, ester or amido,

R<sup>b</sup> is hydrogen, C1-4-alkyl or halogen,

or RaRb is carbonyl.

- (Currently Amended) A compound according to claim 10 wherein R<sup>1</sup> is C3-4-cycloalkyl<sub>5</sub> preferably eyelopropyl.
- (Currently Amended) A compound according to claims 10 wherein R<sup>3</sup> is hydrogen or C1-4-alkyl, preferably hydrogen or methyl.
- 13. (Currently Amended) A compound according to claim 10 wherein

R<sup>a</sup> is hydrogen, methyl, hydroxy, methoxy, methylsulfonyloxy, a Br atom, a F atom or cvano, preferably, hydrogen, methyl, hydroxy or a F atom.

and/or R<sup>b</sup> is hydrogen or methyl, <del>preferably hydrogen,</del> or R<sup>a</sup>R<sup>b</sup> is carbonyl.

(Previously Presented) A compound selected from

1-(6-azetidin-1-vl-2-cvclopropyl-5-methylpyrimidin-4-vl)azepane;

1-[2-cyclopropyl-5-methyl-6-(3-methylazetidin-1-yl)pyrimidin-4-yl]azepane; and pharmaceutically acceptable salts thereof.

- 15. (Previously Presented) A compound according to claim 1 as a pure enantiomer.
- (Previously Presented) A pharmaceutical composition comprising an effective amount of a compound according to claim 1 in combination with a pharmaceutically acceptable diluent or carrier.
- (Original) A pharmaceutical composition according to claim 16 for administration by inhalation
- 18-19. (Canceled)

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20. (Currently Amended) A method for treating respiratory disorders in connection with Chronic Obstructive Pulmonary Disease or for treating symptoms related to chronic bronchitis, emphysema, cough, cystic fibrosis, pulmonary fibrosis, adult respiratory distress syndrome, rhinitis or asthma-comprising administering a therapeutically effective amount of at least one compound according to claim 1 or a pharmaceutically acceptable salt thereof to a patient.

## 21-25. (Canceled)

- (Previously Presented) A pharmaceutical composition comprising an effective amount of a compound according to claim 7 in combination with a pharmaceutically acceptable diluent or carrier.
- (Currently Amended) A pharmaceutical composition comprising an effective amount of a compound according to claim 14 in combination with a pharmaceutically acceptable diluent or emiercarrier.
- 28. (Previously Presented) A compound according to claim 7 as a pure enantiomer.
- 29. (Previously Presented) A compound according to claim 14 as a pure enantiomer.
- (Previously Presented) A pharmaceutical composition according to claim 26 for administration by inhalation.
- (Previously Presented) A pharmaceutical composition according to claim 27 for administration by inhalation.
- 32. (Currently Amended) A method for treating respiratory disorders in connection with Chronic Obstructive Pulmonary Disease or for treating symptoms related to chronic bronchitis, emphysema, cough, cystic fibrosis, pulmonary fibrosis, adult respiratory distress syndrome, rhinitis or asthma-comprising administering a therapeutically effective amount of at least one compound according to claim 7 or a pharmaceutically acceptable salt thereof to a patient.
- (Currently Amended) A method for treating respiratory disorders in connection with Chronic Obstructive Pulmonary Disease or for treating symptoms related to chronic

bronchitis, emphysema, cough, cystic fibrosis, pulmonary fibrosis, adult respiratory distress syndrome, rhinitis or asthma-comprising administering a therapeutically effective amount of at least one compound according to claim 14 or a pharmaceutically acceptable salt thereof to a patient.

- (New) A compound according to claim 4 wherein R<sup>1</sup> is cyclopropyl, isopropyl, cyclobutyl, cyclopentyl, 2-methyl-cyclopropyl or cyclopropylmethyl.
- 35. (New) A compound according to claim 5 wherein R<sup>2</sup> is cyclopropyl or cyclobutyl.
- 36. (New) A compound according to claim 6 wherein R<sup>4</sup> is hydrogen or methyl.
- (New) A compound according to claim 9 wherein NR<sup>4</sup>R<sup>5</sup> is 1-azepanyl, which may be substituted.
- 38. (New) A compound according to claim 11 wherein R<sup>1</sup> is cyclopropyl.
- 39. (New) A compound according to claims 12 wherein R<sup>3</sup> is hydrogen or methyl.
- (New) A compound according to claim 13 wherein R<sup>a</sup> is hydrogen, methyl, hydroxy or a F atom, and/or R<sup>b</sup> is hydrogen.